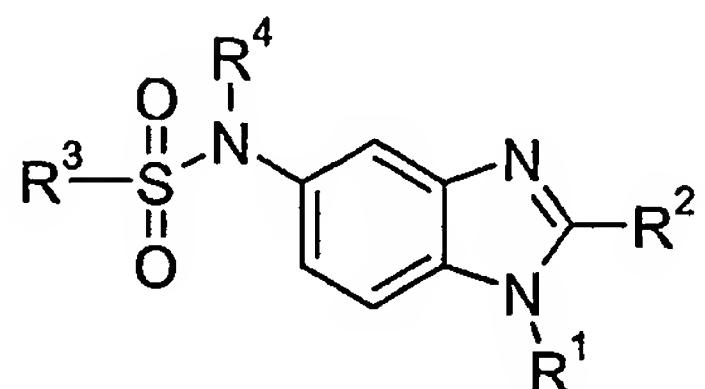


What is claimed is:

1. A compound of Formula I or a pharmaceutically acceptable salt thereof:



5

I

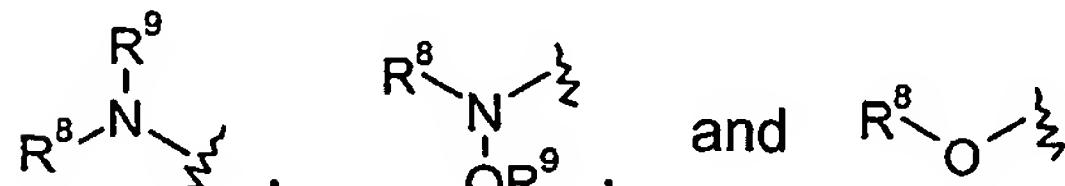
wherein

- R¹ is selected from C₁₋₁₀alkyl, C₂₋₁₀alkenyl, C₂₋₁₀alkynyl, R⁵R⁶N-C₁₋₆alkyl,
R⁵O-C₁₋₆alkyl, R⁵C(=O)N(-R⁶)-C₁₋₆alkyl, R⁵R⁶NS(=O)₂-C₁₋₆alkyl, R⁵CS(=O)₂N(-R⁶)-
C₁₋₆alkyl, R⁵R⁶NC(=O)N(-R⁷)-C₁₋₆alkyl, R⁵R⁶NS(=O)₂N(R⁷)-C₁₋₆alkyl, C₆₋₁₀aryl-
10 C₁₋₆alkyl, C₆₋₁₀aryl-C(=O)-C₁₋₆alkyl, C₃₋₁₀cycloalkyl-C₁₋₆alkyl, C₄₋₈cycloalkenyl-
C₁₋₆alkyl, C₃₋₆heterocyclyl-C₁₋₆alkyl, C₃₋₆heterocyclyl-C(=O)-C₁₋₆alkyl,
C₁₋₁₀hydrocarbyl amino, R⁵R⁶N-, R⁵O-, R⁵C(=O)N(-R⁶)-, R⁵R⁶NS(=O)₂-,
R⁵CS(=O)₂N(-R⁶)-, R⁵R⁶NC(=O)N(-R⁷)-, R⁵R⁶NS(=O)₂N(R⁷)-, C₆₋₁₀aryl, C₆₋₁₀aryl-
C(=O)-, C₃₋₁₀cycloalkyl, C₄₋₈cycloalkenyl, C₃₋₆heterocyclyl and C₃₋₆heterocyclyl-
15 C(=O)-; wherein said C₁₋₁₀alkyl, C₂₋₁₀alkenyl, C₂₋₁₀alkynyl, C₆₋₁₀aryl-C₁₋₆alkyl,
C₆₋₁₀aryl-C(=O)-C₁₋₆alkyl, C₃₋₁₀cycloalkyl-C₁₋₆alkyl, C₄₋₈cycloalkenyl-C₁₋₆alkyl,
C₃₋₆heterocyclyl-C₁₋₆alkyl, C₃₋₆heterocyclyl-C(=O)-C₁₋₆alkyl, C₁₋₁₀hydrocarbyl amino,
C₆₋₁₀aryl, C₆₋₁₀aryl-C(=O)-, C₃₋₁₀cycloalkyl, C₄₋₈cycloalkenyl, C₃₋₆heterocyclyl or
C₃₋₆heterocyclyl-C(=O)- used in defining R¹ is optionally substituted by one or more
20 groups selected from halogen, cyano, nitro, methoxy, ethoxy, methyl, ethyl, hydroxy,
and -NR⁵R⁶;

- R² is selected from C₁₋₁₀alkyl, C₂₋₁₀alkenyl, C₂₋₁₀alkynyl, C₃₋₁₀cycloalkyl, C₃₋₁₀cycloalkyl-C₁₋₆alkyl, C₄₋₈cycloalkenyl-C₁₋₆alkyl, C₃₋₆heterocycloalkyl-C₁₋₆alkyl, C₄₋₈cycloalkenyl, R⁵R⁶N-, C₃₋₅heteroaryl, C₆₋₁₀aryl and C₃₋₆heterocycloalkyl, wherein
25 said C₁₋₁₀alkyl, C₂₋₁₀alkenyl, C₂₋₁₀alkynyl, C₃₋₈cycloalkyl, C₃₋₈cycloalkyl-C₁₋₆alkyl,
C₄₋₈cycloalkenyl-C₁₋₆alkyl, C₃₋₆heterocycloalkyl-C₁₋₆alkyl, C₄₋₈cycloalkenyl, C₃₋₅heteroaryl, C₆₋₁₀aryl or C₃₋₆heterocycloalkyl used in defining R² is optionally
substituted by one or more groups selected from halogen, cyano, nitro, methoxy,
ethoxy, methyl, ethyl, hydroxy, and -NR⁵R⁶;

wherein R⁵, R⁶ and R⁷ are independently selected from -H, C₁₋₆alkyl, C₂₋₆alkenyl, C₂₋₆alkynyl, and a divalent C₁₋₆group that together with another divalent R⁵, R⁶ or R⁷ forms a portion of a ring;

- R³ is selected from -H, C₁₋₁₀alkyl, C₂₋₁₀alkenyl, C₂₋₁₀alkynyl, C₃₋₁₀cycloalkyl,
5 C₃₋₁₀cycloalkyl-C₁₋₆alkyl, C₄₋₈cycloalkenyl-C₁₋₆alkyl, C₃₋₆heterocycloalkyl,


optionally substituted with one or more groups selected from C₁₋₆alkyl, halogen, amino and C₁₋₆alkoxy;

each of R⁸ and R⁹ is independently selected from -H, C₁₋₁₀alkyl, C₂₋₁₀alkenyl, C₂₋₁₀alkynyl, C₃₋₁₀cycloalkyl, C₃₋₁₀cycloalkyl-C₁₋₆alkyl, C₃₋₆heterocyclyl, C₆₋₁₀aryl,

- 10 C₃₋₆heterocyclyl-C₁₋₆alkyl, C₆₋₁₀aryl-C₁₋₆alkyl, and a divalent C₁₋₆group that together with another divalent group selected from R⁸ and R⁹ forms a portion of a ring, wherein said C₁₋₁₀alkyl, C₂₋₁₀alkenyl, C₂₋₁₀alkynyl, C₃₋₁₀cycloalkyl, C₃₋₁₀cycloalkyl-C₁₋₆alkyl, C₃₋₆heterocyclyl, C₆₋₁₀aryl, C₃₋₆heterocyclyl-C₁₋₆alkyl, C₆₋₁₀aryl-C₁₋₆alkyl, or divalent C₁₋₆group is optionally substituted by one or more groups selected from

- 15 halogen, cyano, nitro, methoxy, ethoxy, methyl, ethyl, hydroxy, and -NR⁵R⁶; and

R⁴ is selected from -H, C₁₋₁₀alkyl, C₂₋₁₀alkenyl, C₂₋₁₀alkynyl, C₃₋₁₀cycloalkyl, C₃₋₁₀cycloalkyl-C₁₋₆alkyl, and C₄₋₈cycloalkenyl-C₁₋₆alkyl.

2. A compound as claimed in claim 1, wherein

- 20 R¹ is selected from C₁₋₆alkyl, C₂₋₆alkenyl, C₂₋₆alkynyl, phenyl-C₁₋₄alkyl, C₃₋₁₀cycloalkyl-C₁₋₄alkyl, C₄₋₆cycloalkenyl-C₁₋₄alkyl, C₃₋₁₀heterocyclyl-C₁₋₄alkyl, C₆₋₁₀aryl, C₃₋₁₀cycloalkyl, C₃₋₁₀heterocyclyl and C₄₋₆cycloalkenyl, wherein said C₁₋₆alkyl, C₂₋₆alkenyl, C₂₋₆alkynyl, phenyl-C₁₋₄alkyl, C₃₋₁₀cycloalkyl-C₁₋₄alkyl, C₄₋₆cycloalkenyl-C₁₋₄alkyl, C₃₋₁₀heterocyclyl-C₁₋₄alkyl, C₆₋₁₀aryl, C₃₋₁₀cycloalkyl, C₃₋₁₀heterocyclyl and 25 C₄₋₆cycloalkenyl used in defining R¹ is optionally substituted by one or more groups selected from halogen, cyano, nitro, methoxy, ethoxy, methyl, ethyl, hydroxy, and -NR⁵R⁶;

R² is selected from C₁₋₆alkyl, C₂₋₆alkenyl, C₃₋₆cycloalkyl, C₃₋₆cycloalkyl-C₁₋₄alkyl, C₄₋₆cycloalkenyl-C₁₋₄alkyl, C₃₋₆heterocycloalkyl-C₁₋₄alkyl, C₄₋

- 30 6cycloalkenyl, C₃₋₅heteroaryl, R⁵R⁶N-, and phenyl, wherein said C₁₋₆alkyl, C₂₋₆alkenyl, C₃₋₆cycloalkyl, C₃₋₆cycloalkyl-C₁₋₄alkyl, C₄₋₆cycloalkenyl-C₁₋₄alkyl,

C_{3-6} heterocycloalkyl- C_{1-4} alkyl, C_{4-6} cycloalkenyl, C_{3-5} heteroaryl, R^5R^6N- , and phenyl used in defining R^2 is optionally substituted by one or more groups selected from halogen, cyano, nitro, methoxy, ethoxy, methyl, ethyl, hydroxy and $-NR^5R^6$;

R^3 is selected from $-H$, C_{1-6} alkyl, C_{2-6} alkenyl, C_{3-6} cycloalkyl, C_{3-6}

- 5 $\overset{R^8}{\underset{\diagdown}{N}}\text{---}\overset{R^9}{\underset{\diagup}{N}}$ and $\overset{R^8}{\underset{\diagup}{O}}\text{---}\overset{\diagdown}{\text{Z}}$ optionally substituted with one or
more groups selected from C_{1-6} alkyl and halogen;
each of R^8 and R^9 is independently selected from $-H$, C_{1-6} alkyl, C_{2-6} alkenyl,
 C_{3-6} cycloalkyl, C_{3-6} cycloalkyl- C_{1-6} alkyl, C_{3-6} heterocycl and C_{3-6} heterocycl- C_{1-6} alkyl, wherein said C_{1-6} alkyl, C_{2-6} alkenyl, C_{3-6} cycloalkyl, C_{3-6} cycloalkyl- C_{1-6} alkyl,
10 C_{3-6} heterocycl, C_{3-6} heterocycl- C_{1-6} alkyl and a divalent C_{1-6} group that together
with another divalent group selected from R^8 and R^9 forms a portion of a ring,
wherein said C_{1-6} alkyl, C_{2-6} alkenyl, C_{3-6} cycloalkyl, C_{3-6} cycloalkyl- C_{1-6} alkyl, C_{3-6} heterocycl and C_{3-6} heterocycl- C_{1-6} alkyl, wherein said C_{1-6} alkyl, C_{2-6} alkenyl, C_{3-6} cycloalkyl, C_{3-6} cycloalkyl- C_{1-6} alkyl, C_{3-6} heterocycl, C_{3-6} heterocycl- C_{1-6} alkyl or
15 divalent C_{1-6} group are optionally substituted by one or more groups selected from
halogen, cyano, nitro, methoxy, ethoxy, methyl, ethyl, hydroxy and $-NR^5R^6$; and
 R^4 , R^5 and R^6 are independently selected from $-H$ and C_{1-3} alkyl.

3. A compound as claimed claim 1,

- 20 wherein R^1 is selected from C_{1-6} alkyl, C_{2-6} alkenyl, phenyl- C_{1-4} alkyl,
 C_{3-10} cycloalkyl- C_{1-4} alkyl, C_{4-6} cycloalkenyl- C_{1-4} alkyl, C_{6-10} aryl, C_{3-10} cycloalkyl,
 C_{3-6} heterocycloalkyl- C_{1-4} alkyl, and C_{4-6} cycloalkenyl, wherein said C_{1-6} alkyl, C_{2-6} alkenyl, phenyl- C_{1-4} alkyl, C_{3-10} cycloalkyl- C_{1-4} alkyl, C_{4-6} cycloalkenyl- C_{1-4} alkyl, C_{6-10} aryl, C_{3-10} cycloalkyl, C_{3-6} heterocycloalkyl- C_{1-4} alkyl, and C_{4-6} cycloalkenyl used in
25 defining R^1 is optionally substituted by one or more groups selected from halogen,
methoxy, ethoxy, methyl, ethyl, hydroxy, and $-NR^5R^6$;

- R^2 is selected from C_{1-6} alkyl, C_{2-6} alkenyl, C_{3-6} cycloalkyl and C_{3-6} cycloalkyl- C_{1-4} alkyl, wherein said C_{1-6} alkyl, C_{2-6} alkenyl, C_{3-6} cycloalkyl and C_{3-6} cycloalkyl- C_{1-4} alkyl used in defining R^2 is optionally substituted by one or more groups selected
30 from halogen, methoxy, ethoxy, methyl, ethyl, hydroxy and $-NR^5R^6$;

R^3 is selected from $\text{C}_{2-6}\text{alkyl}$, $\text{C}_{3-6}\text{heterocycloalkyl}$ and $\text{R}^9-\overset{\text{R}^8}{\underset{\text{N}}{\swarrow}}$ optionally substituted with one or more $\text{C}_{1-6}\text{alkyl}$, and;

- wherein said $\text{C}_{3-6}\text{heterocycloalkyl}$ contain at least one nitrogen ring atom and the radical of $\text{C}_{3-6}\text{heterocycloalkyl}$ is located on the at least one nitrogen ring atom,
- 5 and wherein each of R^8 and R^9 is independently selected from $-\text{H}$, $\text{C}_{1-6}\text{alkyl}$, morpholinyl- $\text{C}_{1-3}\text{alkyl}$, pyrrolidinyl- $\text{C}_{1-3}\text{alkyl}$, and piperidinyl- $\text{C}_{1-3}\text{alkyl}$, wherein said $\text{C}_{1-6}\text{alkyl}$, morpholinyl- $\text{C}_{1-3}\text{alkyl}$, pyrrolidinyl- $\text{C}_{1-3}\text{alkyl}$, and piperidinyl- $\text{C}_{1-3}\text{alkyl}$ are optionally substituted by one or more groups selected from halogen, methoxy, ethoxy, methyl, ethyl, hydroxy and $-\text{NR}^5\text{R}^6$; and
10. R^4 , R^5 and R^6 are independently selected from $-\text{H}$ and $\text{C}_{1-3}\text{alkyl}$.

4. A compound as claimed in claim 1, wherein

R^1 is selected from cyclohexylmethyl, cyclopentylmethyl, cyclobutylmethyl, cyclopropylmethyl, 4,4-difluorocyclohexanemethyl, cyclohexylethyl,

15 cyclopentylethyl, tetrahydropyranylmethyl, tetrahydrofuranylmethyl, 1-piperidinylethyl, N-methyl-2-piperidinyl-methyl and benzyl;

R^2 is selected from t-butyl, n-butyl, 2-methyl-2-butyl, isopentyl, 2-methoxy-2-propyl, 2-hydroxy-propyl, trifluoromethyl, 1,1-difluoroethyl, 2,2,2-trifluoroethyl, 1-cyclopropyl-ethyl, 1-methyl-propyl, 1,1-dimethyl-propyl, 1,1-dimethyl-3-buten-1-yl,

20 ethyl, and 2-propyl;

R^3 is $\text{C}_{2-5}\text{alkyl}$ and $\text{R}^8\text{R}^9\text{N}-$, wherein R^8 and R^9 are independently selected from $-\text{H}$, and $\text{C}_{1-3}\text{alkyl}$.

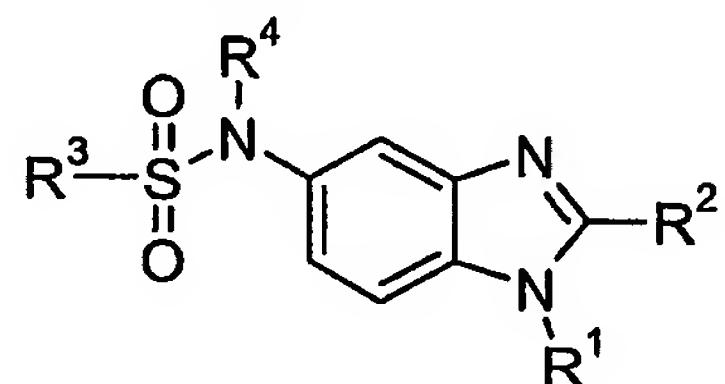
5. A compound selected from:

25 $N-[2\text{-}tert\text{-}Butyl\text{-}1\text{-}(cyclohexylmethyl)\text{-}1H\text{-}benzimidazol\text{-}5\text{-}yl}]\text{-}N,N',N'$ -trimethylsulfamide;

$N-[2\text{-}tert\text{-}Butyl\text{-}1\text{-}(cyclohexylmethyl)\text{-}1H\text{-}benzimidazol\text{-}5\text{-}yl}]\text{-}N',N'\text{-diethyl-N'-methylsulfamide}$;

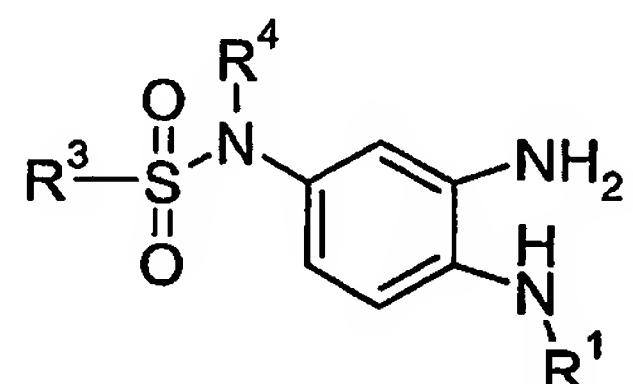
- 5 *N*-[1-(cyclohexylmethyl)-2-(1,1-dimethylpropyl)-1*H*-benzimidazol-5-yl]-*N,N*-dimethyl-sulfamide;
- 10 *N*-[2-*tert*-Butyl-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-yl]-*N*-methylbutane-1-sulfonamide;
- 15 *N*-[2-*tert*-Butyl-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-yl]-*N*-methyl-2-pyrrolidin-1-ylethane sulfonamide;
- 20 *N*-[2-*tert*-Butyl-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-yl]-*N*-methyl-2-morpholin-4-ylethane sulfonamide;
- 25 *N*-[2-*tert*-Butyl-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-yl]-2-[(2-hydroxyethyl)amino]-*N*-methylethane sulfonamide;
- 30 2-(2-Aminoethoxy)-*N*-[2-*tert*-butyl-1-(tetrahydro-2*H*-pyran-4-ylmethyl)-1*H*-benzimidazol-5-yl]-*N*-methylethane sulfonamide;
- N*-{2-*tert*-Butyl-1-[(4,4-difluorocyclohexyl)methyl]-1*H*-benzimidazol-5-yl}-*N*-methylbutane-1-sulfonamide;
- N*-{2-*tert*-Butyl-1-[(4,4-difluorocyclohexyl)methyl]-1*H*-benzimidazol-5-yl}-*N*-methyl-2-piperidin-1-ylethane sulfonamide and pharmaceutically acceptable salts thereof.

6. A compound according to any one of claims 1-5 for use as a medicament.
7. The use of a compound according to any one of claims 1-5 in the manufacture
5 of a medicament for the therapy of pain.
8. The use of a compound according to any one of claims 1-5 in the manufacture
of a medicament for the treatment of anxiety disorders.
- 10 9. The use of a compound according to any one of claims 1-5 in the manufacture
of a medicament for the treatment of cancer, multiple sclerosis, Parkinson's disease,
Huntington's chorea, Alzheimer's disease, gastrointestinal disorders and
cardiovascular disorders.
- 15 10. A pharmaceutical composition comprising a compound according to any one
of claims 1-5 and a pharmaceutically acceptable carrier.
11. A method for the therapy of pain in a warm-blooded animal, comprising the
step of administering to said animal in need of such therapy a therapeutically effective
20 amount of a compound according to any one of claims 1-5.
12. A method for preparing a compound of Formula I,



I

- 25 comprising the step of reacting a compound of Formula II,



II

with a compound of $R^2C(=O)X$, in the presence of a base and optionally a coupling reagent, followed by treatment by an acid;
wherein

- 5 X is selected from Cl, Br, F and OH;
- R^1 is selected from C_{1-10} alkyl, C_{2-10} alkenyl, C_{2-10} alkynyl, $R^5R^6N-C_{1-6}$ alkyl, R^5O-C_{1-6} alkyl, $R^5C(=O)N(-R^6)-C_{1-6}$ alkyl, $R^5R^6NS(=O)_2-C_{1-6}$ alkyl, $R^5CS(=O)_2N(-R^6)-C_{1-6}$ alkyl, $R^5R^6NC(=O)N(-R^7)-C_{1-6}$ alkyl, $R^5R^6NS(=O)_2N(R^7)-C_{1-6}$ alkyl, C_{6-10} aryl- C_{1-6} alkyl, C_{6-10} aryl- $C(=O)-C_{1-6}$ alkyl, C_{3-10} cycloalkyl- C_{1-6} alkyl, C_{4-8} cycloalkenyl-
- 10. C_{1-6} alkyl, C_{3-6} heterocyclyl- C_{1-6} alkyl, C_{3-6} heterocyclyl- $C(=O)-C_{1-6}$ alkyl, C_{1-10} hydrocarbyl amino, R^5R^6N- , R^5O- , $R^5C(=O)N(-R^6)-$, $R^5R^6NS(=O)_2-$, $R^5CS(=O)_2N(-R^6)-$, $R^5R^6NC(=O)N(-R^7)-$, $R^5R^6NS(=O)_2N(R^7)-$, C_{6-10} aryl, C_{6-10} aryl- $C(=O)-$, C_{3-10} cycloalkyl, C_{4-8} cycloalkenyl, C_{3-6} heterocyclyl and C_{3-6} heterocyclyl- $C(=O)-$; wherein said C_{1-10} alkyl, C_{2-10} alkenyl, C_{2-10} alkynyl, C_{6-10} aryl- C_{1-6} alkyl,
- 15. C_{6-10} aryl- $C(=O)-C_{1-6}$ alkyl, C_{3-10} cycloalkyl- C_{1-6} alkyl, C_{4-8} cycloalkenyl- C_{1-6} alkyl, C_{3-6} heterocyclyl- C_{1-6} alkyl, C_{3-6} heterocyclyl- $C(=O)-C_{1-6}$ alkyl, C_{1-10} hydrocarbyl amino, C_{6-10} aryl, C_{6-10} aryl- $C(=O)-$, C_{3-10} cycloalkyl, C_{4-8} cycloalkenyl, C_{3-6} heterocyclyl or C_{3-6} heterocyclyl- $C(=O)-$ used in defining R^1 is optionally substituted by one or more groups selected from halogen, cyano, nitro, methoxy, ethoxy, methyl, ethyl, hydroxy, 20 and $-NR^5R^6$;
- R^2 is selected from C_{1-10} alkyl, C_{2-10} alkenyl, C_{2-10} alkynyl, C_{3-10} cycloalkyl, C_{3-10} cycloalkyl- C_{1-6} alkyl, C_{4-8} cycloalkenyl- C_{1-6} alkyl, C_{3-6} heterocycloalkyl- C_{1-6} alkyl, C_{4-8} cycloalkenyl, R^5R^6N- , C_{3-5} heteroaryl, C_{6-10} aryl and C_{3-6} heterocycloalkyl, wherein said C_{1-10} alkyl, C_{2-10} alkenyl, C_{2-10} alkynyl, C_{3-8} cycloalkyl, C_{3-8} cycloalkyl- C_{1-6} alkyl,
- 25. C_{4-8} cycloalkenyl- C_{1-6} alkyl, C_{3-6} heterocycloalkyl- C_{1-6} alkyl, C_{4-8} cycloalkenyl, C_3 sheteroaryl, C_{6-10} aryl or C_{3-6} heterocycloalkyl used in defining R^2 is optionally substituted by one or more groups selected from halogen, cyano, nitro, methoxy, ethoxy, methyl, ethyl, hydroxy, and $-NR^5R^6$;
- wherein R^5 , R^6 and R^7 are independently selected from -H, C_{1-6} alkyl, C_{2-6} alkenyl, C_{2-6} alkynyl, and a divalent C_{1-6} group that together with another divalent R^5 , R^6 or R^7 forms a portion of a ring;
- R^3 is selected from -H, C_{1-10} alkyl, C_{2-10} alkenyl, C_{2-10} alkynyl, C_{3-10} cycloalkyl, C_{3-10} cycloalkyl- C_{1-6} alkyl, C_{4-8} cycloalkenyl- C_{1-6} alkyl, C_{3-6} heterocycloalkyl,

$\text{R}^8-\overset{\text{R}^9}{\underset{\text{S}}{\text{N}}}-$, $\text{R}^8-\overset{\text{N}}{\underset{\text{OR}^9}{\text{Z}}}$, and $\text{R}^8-\overset{\text{O}}{\underset{\text{Z}}{\text{O}}}$

optionally substituted with one or more

groups selected from C₁₋₆alkyl, halogen, amino and C₁₋₆alkoxy;

each of R⁸ and R⁹ is independently selected from -H, C₁₋₁₀alkyl, C₂₋₁₀alkenyl, C₂₋₁₀alkynyl, C₃₋₁₀cycloalkyl, C₃₋₁₀cycloalkyl-C₁₋₆alkyl, C₃₋₆heterocyclyl, C₆₋₁₀aryl,

- 5 C₃₋₆heterocyclyl-C₁₋₆alkyl, C₆₋₁₀aryl-C₁₋₆alkyl, and a divalent C₁₋₆group that together with another divalent group selected from R⁸ and R⁹ forms a portion of a ring, wherein said C₁₋₁₀alkyl, C₂₋₁₀alkenyl, C₂₋₁₀alkynyl, C₃₋₁₀cycloalkyl, C₃₋₁₀cycloalkyl-C₁₋₆alkyl, C₃₋₆heterocyclyl, C₆₋₁₀aryl, C₃₋₆heterocyclyl-C₁₋₆alkyl, C₆₋₁₀aryl-C₁₋₆alkyl, or divalent C₁₋₆group is optionally substituted by one or more groups selected from

- 10 halogen, cyano, nitro, methoxy, ethoxy, methyl, ethyl, hydroxy, and -NR⁵R⁶; and

R⁴ is selected from -H, C₁₋₁₀alkyl, C₂₋₁₀alkenyl, C₂₋₁₀alkynyl, C₃₋₁₀cycloalkyl, C₃₋₁₀cycloalkyl-C₁₋₆alkyl, and C₄₋₈cycloalkenyl-C₁₋₆alkyl.